

AMENDED IN SENATE AUGUST 17, 1999

AMENDED IN ASSEMBLY JUNE 1, 1999

AMENDED IN ASSEMBLY MAY 6, 1999

AMENDED IN ASSEMBLY MARCH 23, 1999

AMENDED IN ASSEMBLY MARCH 8, 1999

CALIFORNIA LEGISLATURE—1999–2000 REGULAR SESSION

ASSEMBLY BILL

No. 162

Introduced by Assembly Member Runner
(Coauthors: Assembly Members Battin and Cedillo)
(Coauthor: Senator Polanco)

January 15, 1999

An act to amend Section 11100 of the Health and Safety Code, relating to controlled substances.

LEGISLATIVE COUNSEL'S DIGEST

AB 162, as amended, Runner. Controlled substances: ephedrine: retail distributors.

(1) Existing law regulates any manufacturer, wholesaler, retailer, or other person who sells, transfers, or otherwise furnishes specified chemical substances to any person in this state, as specified.

This bill would make it a misdemeanor for any retail distributor to sell in a single transaction more than—4 3 packages of a product that he or she knows to contain ephedrine, pseudoephedrine, norpseudoephedrine, or phenylpropanolamine, or to knowingly sell more than—6 9

grams of ephedrine, pseudoephedrine, norpseudoephedrine, or phenylpropanolamine. The bill ~~also~~ would make an exception for pediatric liquid forms, as defined. The bill would also make the—4 3 packages per transaction limitation or—6 9 grams per transaction limitation applicable to any product that is lawfully sold, transferred, or furnished over the counter without a prescription pursuant to specified provisions of federal law, except as specified. The bill would make clarifying changes. By creating a new crime, this bill would impose a state-mandated local program.

This bill would also provide that it is the intent of the Legislature that specified provisions of state law shall preempt all local ordinances or regulations governing the sale by a retail distributor of over-the-counter products containing ephedrine, pseudoephedrine, norpseudoephedrine, or phenylpropanolamine.

(2) The California Constitution requires the state to reimburse local agencies and school districts for certain costs mandated by the state. Statutory provisions establish procedures for making that reimbursement.

This bill would provide that no reimbursement is required by this act for a specified reason.

Vote: majority. Appropriation: no. Fiscal committee: yes. State-mandated local program: yes.

The people of the State of California do enact as follows:

- 1 SECTION 1. Section 11100 of the Health and Safety
- 2 Code is amended to read:
- 3 11100. (a) Any manufacturer, wholesaler, retailer, or
- 4 other person in this state who sells, transfers, or otherwise
- 5 furnishes any of the following substances to any person or
- 6 business entity in this state or any other state shall submit
- 7 a report to the Department of Justice of all of those
- 8 transactions:
- 9 (1) Phenyl-2-propanone.
- 10 (2) Methylamine.
- 11 (3) Ethylamine.
- 12 (4) D-lysergic acid.
- 13 (5) Ergotamine tartrate.

- 1 (6) Diethyl malonate.
- 2 (7) Malonic acid.
- 3 (8) Ethyl malonate.
- 4 (9) Barbituric acid.
- 5 (10) Piperidine.
- 6 (11) N-acetylanthranilic acid.
- 7 (12) Pyrrolidine.
- 8 (13) Phenylacetic acid.
- 9 (14) Anthranilic acid.
- 10 (15) Morpholine.
- 11 (16) Ephedrine.
- 12 (17) Pseudoephedrine.
- 13 (18) Norpseudoephedrine.
- 14 (19) Phenylpropanolamine.
- 15 (20) Propionic anhydride.
- 16 (21) Isosafrole.
- 17 (22) Safrole.
- 18 (23) Piperonal.
- 19 (24) Thionylchloride.
- 20 (25) Benzyl cyanide.
- 21 (26) Ergonovine maleate.
- 22 (27) N-methylephedrine.
- 23 (28) N-ethylephedrine.
- 24 (29) N-methylpseudoephedrine.
- 25 (30) N-ethylpseudoephedrine.
- 26 (31) Chloroephedrine.
- 27 (32) Chloropseudoephedrine.
- 28 (33) Hydriodic acid.
- 29 (34) Any of the substances listed by the Department
- 30 of Justice in regulations promulgated pursuant to
- 31 subdivision (b).
- 32 (b) The Department of Justice may adopt rules and
- 33 regulations in accordance with Chapter 3.5
- 34 (commencing with Section 11340) of Part 1 of Division 3
- 35 of Title 2 of the Government Code that add substances to
- 36 subdivision (a) if the substance is a precursor to a
- 37 controlled substance and delete substances from
- 38 subdivision (a). However, no regulation adding or
- 39 deleting a substance shall have any effect beyond March

1 1 of the year following the calendar year during which the
2 regulation was adopted.

3 (c) (1) Any manufacturer, wholesaler, retailer, or
4 other person in this state, prior to selling, transferring, or
5 otherwise furnishing any substance specified in
6 subdivision (a) to any person or business entity in this
7 state or any other state, shall require (A) a letter of
8 authorization from that person or business entity that
9 includes the currently valid business license number or
10 federal Drug Enforcement Administration (DEA)
11 registration number, the address of the business, and a full
12 description of how the substance is to be used, and (B)
13 proper identification from the purchaser. The
14 requirement for a full description of how the substance is
15 to be used does not require the person or business entity
16 to reveal their chemical processes that are typically
17 considered trade secrets and proprietary information.

18 (2) For the purposes of this subdivision, “proper
19 identification” for in-state or out-of-state purchasers
20 includes a valid motor vehicle operator’s license or other
21 official and valid state-issued identification of the
22 purchaser, or individual representing the purchasing
23 business entity, which contains a photograph of the
24 purchaser or purchasing individual, and includes the
25 current domicile or mailing address of the purchaser or
26 purchasing individual, other than a post office box
27 number. “Proper identification” also includes the motor
28 vehicle license number of the motor vehicle used by the
29 purchaser or purchasing individual at the time of transfer
30 or the name of the common carrier and the name and
31 valid motor vehicle operator license number of the driver
32 of the common carrier, and the signature of the
33 purchaser, purchasing individual, or driver of the
34 common carrier. The person selling, transferring, or
35 otherwise furnishing any substance specified in
36 subdivision (a) shall affix his or her signature as a witness
37 to the signature and identification of the purchaser,
38 purchasing individual, or driver of the common carrier.

39 (d) Any manufacturer, wholesaler, retailer, or other
40 person in this state who sells, transfers, or otherwise

1 furnishes a substance specified in subdivision (a) to a
2 person or business entity in this state or any other state
3 shall, not less than 21 days prior to delivery of the
4 substance, submit a report of the transaction, which
5 includes the identification information specified in
6 subdivision (c), to the Department of Justice. However,
7 the Department of Justice may authorize the submission
8 of the reports on a monthly basis with respect to repeated,
9 regular transactions between the furnisher and the
10 recipient involving the substance or substances if the
11 Department of Justice determines that the following
12 exist:

13 (1) A pattern of regular supply of the substance or
14 substances exists between the manufacturer, wholesaler,
15 retailer, or other person who sells, transfers, or otherwise
16 furnishes the substance or substances and the recipient of
17 the substance or substances.

18 (2) The recipient has established a record of utilization
19 of the substance or substances for lawful purposes.

20 (e) This section shall not apply to any of the following:

21 (1) Any pharmacist or other authorized person who
22 sells or furnishes a substance upon the prescription of a
23 physician, dentist, podiatrist, or veterinarian.

24 (2) Any physician, dentist, podiatrist, or veterinarian
25 who administers or furnishes a substance to his or her
26 patients.

27 (3) Any manufacturer licensed by the State
28 Department of Health Services or wholesaler licensed by
29 the California State Board of Pharmacy who sells,
30 transfers, or otherwise furnishes a substance to a licensed
31 pharmacy, physician, dentist, podiatrist, veterinarian, or
32 retail distributor as defined in subdivision (h), provided
33 that the manufacturer or wholesaler submits records of
34 any suspicious sales or transfers as determined by the
35 Department of Justice.

36 (4) (A) Any sale, transfer, furnishing, or receipt of
37 any drug which contains ephedrine, pseudoephedrine,
38 norpseudoephedrine, or phenylpropanolamine and
39 which is lawfully sold, transferred, or furnished over the
40 counter without a prescription pursuant to the federal

1 Food, Drug, and Cosmetic Act (21 U.S.C. Sec. 301 et seq.)
2 or regulations adopted thereunder. However, this
3 subparagraph shall apply to preparations in solid or liquid
4 dosage form, *except pediatric liquid forms, as defined,*
5 containing ephedrine, pseudoephedrine,
6 norpseudoephedrine, or phenylpropanolamine ~~as the~~
7 ~~only active medicinal ingredient, or any ephedrine~~
8 ~~combination preparation. In no instance shall the sale of~~
9 ~~any product containing ephedrine, pseudoephedrine,~~
10 ~~norpseudoephedrine, or phenylpropanolamine be~~
11 ~~greater than six grams in a single transaction. where the~~
12 ~~individual transaction involves more than three packages~~
13 ~~or nine grams of ephedrine, pseudoephedrine,~~
14 ~~norpseudoephedrine, or phenylpropanolamine.~~

15 (B) Any ephedrine, pseudoephedrine,
16 norpseudoephedrine, or phenylpropanolamine product
17 subsequently removed from exemption pursuant to
18 Section 814 of Title 21 of the United States Code shall
19 similarly no longer be exempt from any state reporting or
20 permitting requirement, *unless otherwise reinstated*
21 *pursuant to subdivision (b) or (d) of Section 814 of Title*
22 *21 of the United States Code as an exempt product.*

23 (5) Any transfer of a substance specified in subdivision
24 (a) for purposes of lawful disposal as waste.

25 (f) (1) Any person specified in subdivision (a) or (d)
26 who does not submit a report as required by that
27 subdivision or who knowingly submits a report with false
28 or fictitious information shall be punished by
29 imprisonment in a county jail not exceeding six months,
30 by a fine not exceeding five thousand dollars (\$5,000), or
31 by both the fine and imprisonment.

32 (2) Any person specified in subdivision (a) or (d) who
33 has previously been convicted of a violation of paragraph
34 (1) shall, upon a subsequent conviction thereof, be
35 punished by imprisonment in the state prison, or by
36 imprisonment in a county jail not exceeding one year, by
37 a fine not exceeding one hundred thousand dollars
38 (\$100,000), or by both the fine and imprisonment.

39 (g) (1) It is unlawful for any manufacturer,
40 wholesaler, retailer, or other person to sell, transfer, or

1 otherwise furnish a substance specified in subdivision (a)
2 to a person under 18 years of age.

3 (2) It is unlawful for any person under 18 years of age
4 to possess a substance specified in subdivision (a).

5 (3) Notwithstanding any other law, it is unlawful for
6 any retail distributor to (i) sell in a single transaction
7 more than ~~four~~ *three* packages of a product that he or she
8 knows to contain ephedrine, pseudoephedrine,
9 norpseudoephedrine, or phenylpropanolamine, or (ii)
10 knowingly sell more than ~~six~~ *nine* grams of ephedrine,
11 pseudoephedrine, norpseudoephedrine, or
12 phenylpropanolamine, *other than pediatric liquids as*
13 *defined*. Except as otherwise provided in this section, the
14 ~~four~~ *three* package per transaction limitation or ~~six~~ *nine*
15 gram per transaction limitation imposed by this
16 paragraph shall apply to any product that is lawfully sold,
17 transferred, or furnished over the counter without a
18 prescription pursuant to the federal Food, Drug, and
19 Cosmetic Act (21 U.S.C. Sec. 301 et seq.), or regulations
20 adopted thereunder, ~~and that has been~~ *unless* exempted
21 from the requirements of the federal Controlled
22 Substances Act by the federal Drug Enforcement
23 Administration pursuant to Section 814 of Title 21 of the
24 United States Code.

25 (4) A violation of this subdivision is a misdemeanor.

26 (h) For the purposes of this article, the following terms
27 have the following meanings:

28 (1) “Drugstore” is any entity described in Code 5912
29 of the Standard Industrial Classification (SIC) Manual
30 published by the United States Office of Management
31 and Budget, 1987 edition.

32 (2) “General merchandise store” is any entity
33 described in Codes 5311 to 5399, inclusive, and Code 5499
34 of the Standard Industrial Classification (SIC) Manual
35 published by the United States Office of Management
36 and Budget, 1987 edition.

37 (3) “Grocery store” is any entity described in Code
38 5411 of the Standard Industrial Classification (SIC)
39 Manual published by the United States Office of
40 Management and Budget, 1987 edition.

1 (4) “Ordinary over-the-counter ephedrine,
2 pseudoephedrine, norpseudoephedrine, or
3 phenylpropanolamine product” means a product
4 containing ephedrine, pseudoephedrine,
5 norpseudoephedrine, or phenylpropanolamine sold in
6 package sizes of not more than 3.0 grams of ephedrine,
7 pseudoephedrine, norpseudoephedrine, or
8 phenylpropanolamine, and is packaged in blister packs,
9 each blister containing not more than two dosage units,
10 or where the use of blister packs is technically infeasible,
11 is packaged in unit dose packets or pouches; or, if a liquid,
12 sold in package sizes of not more than 3.0 grams of
13 ephedrine, pseudoephedrine, norpseudoephedrine, or
14 phenylpropanolamine.

15 (5) “*Pediatric liquid forms*” means _____.

16 (6) “Retail distributor” means a grocery store, general
17 merchandise store, drugstore, or other related entity, the
18 activities of which, as a distributor of ephedrine,
19 pseudoephedrine, norpseudoephedrine, or
20 phenylpropanolamine products, are limited exclusively
21 to the sale of ephedrine, pseudoephedrine,
22 norpseudoephedrine, or phenylpropanolamine products
23 for personal use both in number of sales and volume of
24 sales, either directly to walk-in customers or in
25 face-to-face transactions by direct sales. “Retail
26 distributor” includes an entity that makes a direct sale,
27 but does not include the parent company of that entity if
28 the company is not involved in direct sales regulated by
29 this article.

30 ~~(6)~~—

31 (7) “Sale for personal use” means the sale in a single
32 transaction to an individual customer for a legitimate
33 medical use of a product containing ephedrine,
34 pseudoephedrine, norpseudoephedrine, or
35 phenylpropanolamine in dosages at or below that
36 specified in paragraph (3) of subdivision (g). “Sale for
37 personal use” also includes the sale of those products to
38 employers to be dispensed to employees from first-aid
39 kits or medicine chests.

1 (i) It is the intent of the Legislature that this section
2 shall preempt all local ordinances or regulations
3 governing the sale by a retail distributor of
4 over-the-counter products containing ephedrine,
5 pseudoephedrine, norpseudoephedrine, or
6 phenylpropanolamine.

7 SEC. 2. No reimbursement is required by this act
8 pursuant to Section 6 of Article XIII B of the California
9 Constitution because the only costs that may be incurred
10 by a local agency or school district will be incurred
11 because this act creates a new crime or infraction,
12 eliminates a crime or infraction, or changes the penalty
13 for a crime or infraction, within the meaning of Section
14 17556 of the Government Code, or changes the definition
15 of a crime within the meaning of Section 6 of Article
16 XIII B of the California Constitution.

